- TI Design of synthetic hexapeptide ***substrates*** for prostate-specific antigen using single-position minilibraries
- AU Yang, C. F.; Porter, E. S.; Boths, J.; Kanyi, D.; Hsieh, M.; Cooperman, B. S.
- CS Department of Chemistry, Rowan University, Glassboro, NJ, 08028, USA
- SO Journal of Peptide Research (1999), 54(5), 444-448 CODEN: JPERFA; ISSN: 1397-002X
- PB Munksgaard International Publishers Ltd.
- DT Journal
- LA English
- AB Prostate-specific antigen (PSA), a serine endoprotease with chymotrypsin-like ***substrate*** specificity, is a marker used widely for detection of prostate cancer and other prostate diseases, catalyzing hydrolysis of the gel-forming proteins semenogelins I and II, which are synthesized and secreted by the seminal vesicle. In this study we report the use of two single-position minilibraries and RP-HPLC selection to optimize a hexapeptide ***substrate*** for PSA, spanning
 - ***substrate*** positions P3 to P3'. PSA has been shown previously to prefer tyrosine in position ***P1*** (Denmeade, S.R., et al., 1997). Here we demonstrate preference for serine in position ***P1*** ' and strong preference for phenylalanine in position P2. Based on these results we have designed and demonstrated the utility of the optimized fluorogenic PSA ***substrate*** 7-methoxy-coumarin-4-acetylGlnPheTyrSerSerAsnLys(.epsilon.-2,4-dinitrophenyl)amide, which permits continuous monitoring of PSA endopeptidase activity at high sensitivity.
- RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1998:556406 CAPLUS

DN 129:312624

- TI ***Substrate*** specificity of non-pepsin-type acid proteinase, Aspergillus niger proteinase A
- AU Komatsu, Shinji; Nishii, Wataru; Sasaki, Hiroshi; Muramatsu, Tomonari; Tanokura, Masaru
- CS Biotechnology Research Center, University of Tokyo, Tokyo, 113, Japan
- SO Advances in Experimental Medicine and Biology (1998), 436(Aspartic Proteinases), 345-348
 CODEN: AEMBAP; ISSN: 0065-2598
- PB Plenum Publishing Corp.
- DT Journal
- LA English
- AB The peptide ***library*** method was applied to investigate the

 Pl and P2 site specificities of Aspergillus niger proteinase A.

 The resp. peptide ***substrates*** were RGFFXTPRA and RGFXYTPRA. The
 specificity at the ***Pl*** site is higher than that at P2 and is
 considerably complicated. Methionine, phenylalanine and histidine are the
 preferred ***Pl*** site residues. The results on P2 site specificity
 showed that the ***enzyme*** tends to ***cleave*** only those
 peptides which a contain large residue at P2; residue charge did not
 affect specificity.
- RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d his

(FILE 'HOME' ENTERED AT 17:50:58 ON 10 AUG 2005)

FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 17:51:13 ON 10 AUG 2005 42 S HYDROPATHIC INDEX

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ъ3
             0 S HYDROPATHIC INDEX/TI
           798 S HYDROPATHIC
L4
           130 S HYDROPATHIC/TI
L<sub>5</sub>
L6
             3 S L5 AND REVIEW
L7
             3 DUP REM L6 (0 DUPLICATES REMOVED)
L8
           301 S L4 AND PREDICT?
L9
             2 S L8 AND REVIEW
L10
           304 S HYDROPATH?/TI
L11
           174 S L10 NOT L5
L12
             0 S LL1 AND ENZYM? AND SUBSTRAT?
             7 S L11 AND ENZYM?
L13
    FILE 'STNGUIDE' ENTERED AT 17:59:06 ON 10 AUG 2005
     FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 18:00:23 ON 10 AUG 2005
           124 S HYDROPATH? AND ENZYM? AND SUBSTRAT?
L14
L15
             0 S L14 AND REVIEW
L16
            42 S L14 AND (PREDICT? OR UNPREDICT?)
             0 S L14 AND UNPREDICT?
L17
L18
             1 S L16 AND CLEAV?
L19
           16 S L14 AND LIBRARY
L20
             8 DUP REM L19 (8 DUPLICATES REMOVED)
L21
             5 S L14 AND SUBSTIT?
             3 DUP REM L21 (2 DUPLICATES REMOVED)
L22
           55 S HYDROPATH? AND PEPTID? AND MIM?
L23
L24
            22 DUP REM L23 (33 DUPLICATES REMOVED)
L25
            2 S L24 AND REVIEW
L26
             2 S L24 AND ENZYM?
L27
          4559 S HYDROPATH?
L28
            45 S L27 AND OLIGOPEP?
            33 DUP REM L28 (12 DUPLICATES REMOVED)
L29
             0 S L29 AND REVIEW
L30
             5 S L29 AND ENZYM?
L31
L32
            11 S L10 AND REVIEW
             8 S L32 NOT L6
L33
L34
             6 DUP REM L33 (2 DUPLICATES REMOVED)
     FILE 'STNGUIDE' ENTERED AT 18:22:44 ON 10 AUG 2005
     FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 18:24:17 ON 10 AUG 2005
T.35
           103 S L27 AND REVIEW
L36
             5 S L35 AND SUBSTRAT?
L37
             5 DUP REM L36 (0 DUPLICATES REMOVED)
L38
            62 S L27 AND ((OLIGOPEP?) OR (PEP? MIM?) OR PEPTO?)
L39
            39 DUP REM L38 (23 DUPLICATES REMOVED)
L40
             9 S L39 AND (ENZYM? OR SUBSTRAT?)
L41
             9 DUP REM L40 (0 DUPLICATES REMOVED)
          1095 S L27 AND ENZYM?
L42
            13 S L27 AND CONSERV? SUB?
L43
L44
             5 DUP REM L43 (8 DUPLICATES REMOVED)
L45
            24 S L27 AND SUBSTRAT?/TI
L46
             8 DUP REM L45 (16 DUPLICATES REMOVED)
L47
          15695 S SITE DIRECT? AND MUTAGEN? AND SUBSTRAT?
L48
          2511 S L47 AND SUBSTRAT?/TI
L49
            97 S L48 AND CONSERVATI?
L50
             0 S L49 AND REVIEW
L51
            53 DUP REM L49 (44 DUPLICATES REMOVED)
L52
           862 S SUBSTRAT?/TI AND ENZYM? AND LIBRAR?
L53
           233 S L52 AND (CLEAV? OR SISS?)
L54
             0 S L53 AND SISS?
L55
             0 S L53 AND SISC?
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L2

1 S L1 AND REVIEW

L56	241	S	L52	AND	(CLEAV?	OR SC	IS?)				
L5.7	2	S	L56	AND	REVIEW						
L58	2	DU	JP RE	EM L	57 (O DU	PLICAT	ES REMOVEI)			
L59	3	S	L56	AND	HYDROPA	TH?					
L60	981	S	SUBS	STRAT	r? And e	:NZYM?	AND LIBRAF	R? AND	(CLEAV?	OR	SCIS?)
L61	976	S	L60	NOT	(L57 OR	L58 O	R L59)				
L62	15	S	L61	AND	REVIEW						
L63	11	DU	JP RF	EM Le	52 (4 DU	PLICAT	ES REMOVEI))			
L64	130	s	L61	AND	P1						
L65	70	DU	JP RE	EM Le	54 (60 D	UPLICA	TES REMOVE	ED)			
L66	28	S	L65	AND	(ALZH?	OR APP	? OR AMYLO)ID?)			
L67					-		OR AMYLO	[D?)			
L68	8	S	L65	AND	STRUCTU	RE ACT	IVITY				
L69	0	S	L65	AND	REVIEW						
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